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# Runoff- and erosion-driven transport of cattle slurry: linking molecular tracers to hydrological processes

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## Abstract

The addition of cattle slurry to agricultural land is a widespread practise, but if not correctly managed it can pose a contamination risk to aquatic ecosystems. The transport of inorganic and organic components of cattle slurry to watercourses is a major concern, yet little is known about the physical transport mechanisms and associated fluxes and timings of contamination threats. Therefore, the aim of the study was to ascertain the importance of flow pathway partitioning in the transport (fluxes and timing) of dissolved and particulate slurry-derived compounds with implications for off-site contamination. A series of rainfall–runoff and erosion experiments were carried out using the TRACE (Test Rig for Advancing Connectivity Experiments) experimental hillslope facility. The experiments allowed the quantification of the impact of changing slope gradient and rainfall intensity on nutrient transport from cattle slurry applied to the hillslope, via surface, subsurface and vertical percolated flow pathways, as well as particulate transport from erosion. The dissolved components were traced using a combination of ammonium ( $\text{NH}_4^+$ ) and fluorescence analysis, while the particulate fraction was traced using organic biomarkers,  $5\beta$ -stanols. Results showed that rainfall events which produced flashy hydrological responses, resulting in large quantities of surface runoff, were likely to move sediment and also flush dissolved components of slurry-derived material from the slope, increasing the contamination risk. Rainfall events which produced slower hydrological responses were dominated by vertical percolated flows removing less sediment-associated material, but produced leachate which could contaminate deeper soil layers, and potentially groundwater, over a more prolonged period. Overall, this research provides new insights into the partitioning of slurry-derived material when applied to an unvegetated slope and the transport mechanisms by which contamination risks are created.

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## 1 Introduction

Contamination of water bodies due to poor agricultural management is a global problem that affects aquatic ecosystems from individual catchments through to estuaries and oceans, and impacts human water supplies (e.g. Mitsch et al., 2001; Diaz and Rosenberg, 2008; Osterman et al., 2009; HELCOM, 2009). The ongoing introduction of legislation aimed at improving aquatic ecosystems (e.g. European Union Water Framework Directive) has focussed attention on efforts to decrease inputs of pollutants into surface and groundwater. Transport of inorganic and organic nutrients from livestock slurries to water courses are a particular concern, typically associated with over-application or poorly timed application of organic fertilisers to agricultural land (Dungait et al., 2012; Chadwick and Chen, 2003). The inorganic component of slurries (mainly nitrogen, N) is generally considered as more soluble and directly bioavailable compared with the organic fraction, and specific forms such as nitrate are also more liable to immediate loss by leaching into watercourses (e.g. Chambers et al., 2000; Delconte et al., 2014; Quemada et al., 2013). However, studies have shown that the organic fraction can provide a large proportion of the total N load in some catchments and a smaller yet environmentally significant fraction in N-enriched waters (Durand et al., 2011; Willett et al., 2004; Johnes and Burt, 1991). This excess N leads to eutrophication of surface waters, resulting in algal blooms and reduced oxygen levels. Sutton (2011) reported that excess N in the environment costs the EU between EUR 70–320 billion per annum, more than double the value that the N fertilisers provide to EU farms in terms of production. Combating and mitigating against these problems is extremely costly; the UK spends up to GBP 300 million each year cleaning water courses, equating to up to 2 % of the gross agricultural output (Pretty et al., 2000). Given these severe environmental and financial consequences it is important to better understand the main transport pathways, fluxes and transit times of pollutants from livestock slurries into watercourses to develop effective mitigation strategies. Slurry transport has been quantified mainly in terms of the inorganic fraction (e.g. nitrate and phosphate)

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and associated pathogens (e.g. coliforms) (e.g. Edwards et al., 2012; Coelho et al., 2012; Eastman et al., 2010), but there is a relative lack of understanding of physical transport mechanisms and the associated fluxes and timings of contamination threats from areas treated with livestock slurries.

The transport of slurry components on land is primarily controlled by the hydrological and erosion regime operating at the site of application. The dynamics of flow pathway partitioning during storm events and the consequential output of agricultural contaminants has been explored (Delpla et al., 2011; Blanchard and Lerch, 2000; Gao et al., 2004; e.g. Zhang et al., 1997; Malone et al., 2004), but there is a lack of experimental data quantifying contaminant export via individual flow pathways due to methodological challenges. In a field context it is challenging to monitor multiple flow pathways without destructive sampling, although new research platforms are now making this type of research more possible (see Peukert et al., 2014 for an example). There are also challenges when choosing field sites that represent transport regimes across different slope angles or to account for other environmental variables. Therefore, laboratory flume experiments have been widely used to investigate questions relating to hydrology and erosion in conjunction with solute and sediment transport, although most studies to date have only measured surface runoff and/or vertical drainage (e.g. Montenegro et al., 2013; Guo et al., 2010; Aksoy et al., 2012; Asam et al., 2012). Our previous work using a one-dimensional soil column revealed the rapid partitioning of livestock slurries in the soil-water system into sediment-associated material remaining on or close to the surface and dissolved components which moved rapidly through the soil by leaching (Lloyd et al., 2012). Given this partitioning into surface and subsurface components and its potential importance for contamination of downstream aquatic and soil environments, we aim to quantify the relative fluxes of different slurry compounds driven by surface (overland flow and erosion) and subsurface (throughflow and leaching) flow pathways during a series of experiments in which rainfall rate and slope angle vary. We vary rainfall rate and slope angle in order to simulate hydrological variations which may result in differential partitioning of flow pathways.

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surface throughflow and (3) vertical percolated flow, in order to investigate the transport of slurry-derived compounds. Eroded sediment was separated from the surface runoff using the protocol outlined below and kept for analysis. At the end of the rainfall simulation soil cores were taken from the slope according to the sampling strategy outlined in Sect. 2.2. The soil cores and eroded sediment were used to investigate the erosion and within-slope deposition of slurry-derived particulates.

## 2.2 Sampling

Discharge from each flow pathway was determined at regular intervals during the course of the rainfall simulations. Discharge from the surface runoff pathway was monitored by logging water depth using a “V”-notch weir and capacitance depth probe, at 1 min intervals. The water depth was converted to discharge by a pre-calibrated stage–discharge relationship for the “V”-notch weir. Additional manual samples of the surface runoff were taken at ~ 5 min intervals to determine sediment concentration and to obtain eroded sediment samples for analysis. The water and sediment were separated using a centrifuge (2400 rpm, 30 min) and the water samples were filtered through 0.45 µm filter (Whatman, cellulose acetate) then stored at –18 °C until analysis was carried out. Flow discharge from the subsurface and percolated pathways was monitored manually at ~ 5 min intervals by timing the flow of a known volume of water.

At the end of the rainfall simulations, 18 soil cores were taken over a regularly spaced 3 × 6 sample grid leaving a buffer of 0.5 m around the slope perimeter to avoid edge effects (samples 0.75 m apart across slope, 1 m apart downslope), to a depth of 5 cm. Only the top 5 cm were sampled because pilot experiments and previous research has shown that this section is the most important in terms of the transport of slurry-derived material (see Lloyd et al., 2012). The soil cores and eroded sediment were freeze-dried and stored at –18 °C until analysis.

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2.3 Laboratory analyses

2.3.1 Runoff water analyses

All water samples were analysed for  $\text{NH}_4^+$  ( $\mu\text{g L}^{-1}$ ) using a continuous segmented flow autoanalyser (AA3; Seal Analytical).  $\text{NH}_4^+$  was determined using the Berthelot reaction (Berthelot, 1859), where a blue-green compound was produced and quantified colorimetrically.  $\text{NH}_4^+$  was the only inorganic N fraction measured as the cattle slurry used contained no measureable  $\text{NO}_3^-$  or  $\text{NO}_2^-$  (see Table 1). Also, previous work carried out using the same soil and slurry showed that after slurry application and leaching for 8 h, there was no detectable difference between the concentrations of extractable  $\text{NO}_3^-$  or  $\text{NO}_2^-$  from slurry-treated or control soils (unpublished data). As a result, there is no evidence to suggest that nitrification would occur over the time-scales of the current TRACE experiments and therefore  $\text{NH}_4^+$  was the sole inorganic compound used to trace slurry in this paper.

Fluorescence spectroscopic analysis of all leachate samples was carried out using a HORIBA Jobin Yvon FluoroLog®-3 spectrofluorometer (Stanmore, UK); the excitation source was a Xe lamp. Samples were measured in a 1 cm quartz cuvette at room temperature. The methodology of Peuravuori et al. (2002) was adopted to collect synchronous spectra, using 1 nm increments from 250–600 nm, with an 18 nm offset between the excitation and emission monochromators. The spectra were then corrected for the fluorescence of Milli-Q water and for inner-filter effects, based on the sample absorbance. The sample absorbance at 250–600 nm was measured in a 1 cm cuvette, using a Shimadzu UVmini-1240. Milli-Q subtracted absorbance spectra were then used to correct the fluorescence spectra for primary and secondary inner-filter effects using the equation:

$$I_c = I_m / \left( 10^{-b(A_{\text{ex}} + A_{\text{em}})} \right)$$

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where  $I_c$  is the true fluorescence intensity,  $I_m$  is the measured fluorescence intensity,  $b$  is the sample path length and  $A_{ex}$  and  $A_{em}$  are the absorbance values at the excitation and emission wavelengths respectively (Lakowicz, 1983; Ohno, 2002). The spectra were then converted to arbitrary Raman units by normalising the fluorescence intensities by the area of the water Raman peak in order to compare the relative magnitude of the peaks present between the experiments. The ratio between the fluorescence intensities of emissions at  $\sim 290$  nm and  $\sim 380$  nm was calculated as previous work has shown that it can be used to monitor the presence of slurry-derived compounds in natural waters (Baker, 2002b, a; Naden et al., 2010; Lloyd et al., 2012).

### 2.3.2 Soil and eroded sediment analyses

The soil and eroded sediment samples were analysed in triplicate for total carbon (TC), inorganic carbon (IC) and total nitrogen (TN) (%) using a Carlo Erba EA1108 Elemental Analyser. Total organic carbon (TOC) was then calculated by subtracting the IC from the TC.

Lipid analysis was used to extract and quantify the stanol concentration within the soil cores and eroded sediment samples, specifically the  $5\beta$ -stanols, which were used as a tracer of slurry-derived particulates (Bull et al., 2002; Leeming et al., 1996; Evershed et al., 1997; Nash et al., 2005; Lloyd et al., 2012). As  $5\beta$ -stanols stigmastanol and its epimer are the biohydrogenation products of sitosterol (a major plant sterol) and are only produced during rumination, they can be used as unequivocal tracers of the hydrophobic fraction of ruminant faeces. The  $5\alpha$  form is produced by microbial activity under aerobic conditions, i.e. outside the rumen, so the ratio  $5\beta : 5\alpha$  is used to investigate the contribution of either source. To achieve this, a total lipid extract (TLE) was obtained from the soil and eroded sediment samples using Soxhlet extraction ( $\text{CH}_2\text{Cl}_2 : \text{Me}_2\text{CO}$  9 : 1 v/v, 24 h, van Bergen et al., 1997) with an internal standard (preg-5-en-3 $\beta$ -ol). Aliquots of the TLE samples were saponified by adding 1 mL 0.5 M methylated MeOH and heating at 70 °C for 60 min before acidification to pH 3 by adding 1 mL of 1 M HCl. Then 1 mL of DCM-extracted DDW was added along with 2 mL of DCM and the sample

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was vortex mixed for 20 s and allowed to settle to give a two-phase sample. The organic layer was extracted from the bottom then the DCM extraction was repeated two more times to ensure the entire organic sample was collected. The composite organic phase was then filtered through a pipette containing anhydrous sodium sulphate ( $\text{NaSO}_4$ ) in order to ensure any residual water was removed, then the sample was blown down to dryness under  $\text{N}_2$ . An aliquot of the saponified TLE was then derivatised using 50  $\mu\text{L}$  of N,O-bis(trimethylsilyl)trifluoroacetamide with 1 % trimethylchlorosilane and heating at 70 °C for 1 h. The samples were then analysed using a Finnigan TRACE GC/MS by injecting 1  $\mu\text{L}$  on-column, using a HP-1 (50 m, 0.32 mm, 0.17  $\mu\text{m}$ ) column. The GC temperature programme was: 50 °C (2 min), ramping to 245 °C at a rate of 15 °C  $\text{min}^{-1}$ , then increasing to 250 °C at a rate of 0.5 °C  $\text{min}^{-1}$  followed by a ramp to 300 °C at a rate of 6 °C  $\text{min}^{-1}$ , then finally held for 20 min. This GC temperature programme was chosen to maximise the information gained at the time period where the stanols elute. The mass spectrometer was operated at 70 eV with the quadrapole mass analyser scanning the range  $m/z$  50–650 (scan time 0.6 s). The distribution of sterols and stanols present were then quantified against the internal standard.

The ratio of  $5\beta$ - :  $5\alpha$ -stanols was determined in order to assess the relative contribution of slurry-derived insoluble organic matter (IOM) to native soil IOM in the soil cores. This is a useful technique where the  $5\beta$ -stanol concentrations are low, as the ratio allows differences between the samples to be seen more clearly.

## 2.4 Statistical analyses

Statistical differences between the slurry treated experiments and the control experiments were tested for each of the measured parameters using either a  $t$  test or a Mann–Whitney test depending on whether the specific data sets were normally or non-normally distributed.

3 Results

3.1 Flow pathway partitioning and erosion

The results show that as rainfall intensity and slope gradient increased a higher proportion of flow was transported via surface and shallow subsurface flows in the slurry treated experiments (Fig. 2). Doubling the rainfall intensity had a larger impact on the flow pathway partitioning compared with doubling the slope gradient. Figure 2 summarises the partitioning of water flow through the slope for the slurry-treated experiments. Discharge data for the control experiments are not included as the results mirrored the slurry experiments run under the same experimental conditions. The two experimental end-member scenarios showed a reversal in behaviour of the flow partitioning, where at 5°, 60 mm h<sup>-1</sup>, ~ 99 % of the flow was transported as vertical percolated flow and at 10°, 120 mm h<sup>-1</sup>, ~ 99 % of the flow was routed across the soil surface as overland flow. All experiments received the same volume of rainfall, therefore halving the slope gradient and rainfall intensity caused a 60 % increase in the storage of water within the soil. At a rainfall intensity of 60 mm h<sup>-1</sup> doubling the slope gradient induced a reduction in the vertical percolated flow (to 26 %) and increase in subsurface throughflow (28 %) and surface runoff (46 %) due to higher surface water velocities and reduced infiltration rates. Therefore, overland flow generation was a combination of infiltration-excess (fast) and saturation excess (slower) overland flow. When the rainfall intensity increased to 120 mm h<sup>-1</sup> the system was dominated by infiltration-excess overland flow. At a slope gradient of 5°, 67 % of the discharge was surface runoff and 29 % was subsurface throughflow. However, increasing the slope gradient to 10° resulted in 99 % of the flow transported as surface runoff.

Figure 3 shows the corresponding sedigraphs of eroded sediment transported in the overland flow for each of the slurry-treated experiments. Like discharge the data for the control experiments is not included as their behaviour mirrored that of the slurry-treated experiment run under the same experimental conditions. The experiments run at 60 mm h<sup>-1</sup> rainfall produced sediment concentrations that increased gradually until

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the middle of the experiment before decreasing later in the event. At  $120 \text{ mm h}^{-1}$  the  $5^\circ$  experiment produced a similar trend to those run at the lower rainfall intensity (at both  $5^\circ$  and  $10^\circ$  slopes), however the sediment discharges were generally higher. When the slope gradient increased to  $10^\circ$  the shape of the sedigraph changed and the greatest sediment concentrations were at the beginning of the rainfall simulations. This suggests that rainsplash, combined with higher water velocities and higher flow shear stress expected at steeper gradients, could initially rapidly move a large quantity of sediment, after which the system slowed to a sediment export rate similar to that of the lower gradient slope.

### 3.2 Dissolved chemical species analyses

#### 3.2.1 Ammonium ( $\text{NH}_4^+$ )

Figures 4–6 show a direct comparison of the flow and sediment and the  $\text{NH}_4^+$  concentration from each monitored flow pathway during each experiment. The control experiments (Fig. 4) produced a total of between 12 and 25 mg of leached  $\text{NH}_4^+$  originating from the soil (no added slurry), with concentrations never exceeding  $0.1 \text{ mg L}^{-1}$  from any pathway. The total mass of  $\text{NH}_4^+$  exported from each experiment depended on the relative flow pathway partitioning because the experiments which generated higher volumes of surface runoff contributed a larger flux of  $\text{NH}_4^+$ , between 1.8 and 11.3 % of the total  $\text{NH}_4^+$  measured from the slope outlets. An approximately equal proportion of the exported  $\text{NH}_4^+$  was leached from the vertical percolated flow outlets in all control experiments, between 10.9 mg (48 %) 7.6 mg (62 %), with the remainder transported via sub-surface runoff.

The addition of slurry to the slope caused a marked increase in the export of  $\text{NH}_4^+$  compared to the controls, in some cases by an order of magnitude, indicating that the main source of  $\text{NH}_4^+$  leached from the slope was the added slurry. The  $\text{NH}_4^+$  concentrations from the slurry-treated experiments were significantly higher than those from

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the control experiments (Mann–Whitney, control vs. slurry,  $p = < 0.001$ ). As with the control experiments, total  $\text{NH}_4^+$  export was affected by the flow pathway partitioning. The slurry-treated experiments which were dominated by surface runoff exhibited the largest export of  $\text{NH}_4^+$ , between 66 and 99 % of the total. The experiment with the most surface runoff ( $10^\circ$ ,  $120 \text{ mm h}^{-1}$ ) generated a  $\text{NH}_4^+$  yield of 404 mg, an order of magnitude larger than any other experiment.  $\text{NH}_4^+$  was also observed at similar concentrations in the sub-surface flow pathway when activated. During the experiment where the majority of the water flow was via vertically percolated pathways ( $5^\circ$ ,  $60 \text{ mm h}^{-1}$ ), the concentrations of  $\text{NH}_4^+$  were similar to those observed in the dominant pathways for other slurry experiments ( $0.1\text{--}0.5 \text{ mg L}^{-1}$ ). The area associated with the largest percolated flux tended to be the top of the slope directly below the slurry application area, illustrating that the slurry can still be rapidly transported during smaller rainfall events but the dissolved fraction is infiltrated rather than routed across the soil surface.

In general, the transport of  $\text{NH}_4^+$  was controlled by the flow partitioning, and the concentrations did not increase with contact time with the soil, providing additional evidence that the  $\text{NH}_4^+$  was representing the dissolved slurry-derived material. This can be seen very clearly in Fig. 6c, where the concentration of  $\text{NH}_4^+$  observed in the percolated flow pathway is highest after 20 min, showing that the slurry material had reached the bottom of the slope.

### 3.2.2 Fluorescence spectroscopy

Figure 7 shows the average fluorescence spectra for each flow pathway for each experiment, along with the mean spectra of the control samples and the slurry. This plot highlights the difference between the fluorescence spectra of the slurry and control soil is the magnitude of the peak at excitation wavelength 270 nm (emission 290 nm), which has been attributed to “proteinaceous material” (Baker, 2002b, a; Peuravuori et al., 2002), therefore providing a useful tracer of slurry-derived dissolved organic matter (DOM) (Lloyd et al., 2012). This “protein” fluorescence can be a useful tracer









were greater than those in the control soil in only one experiment, however they were not significantly more (Mann–Whitney, control vs. slurry,  $p = 0.179$ ). Overall, these data confirm that the elemental analyses alone do not provide a robust method to trace the transport of slurry-derived particulates.

### 3.3.2 Lipid analyses on eroded sediment and soil cores

Figure 9 shows example partial chromatograms for a soil core from the top area of the slope, eroded sediment and the control soil. These data clearly illustrate the dominance of the  $5\beta$ -stanols in the slurry-treated samples relative to the control soil. The concentration of  $5\beta$ -stanols, a robust biomarker for slurry-derived IOM, in the slurry was  $12\,982\text{ ng g}^{-1}$  compared with an average of  $67.7\text{ ng g}^{-1}$  in the control soil. The soil cores taken from the slurry application area were significantly enriched with concentrations averaging  $2850\text{ ng g}^{-1}$  (Mann–Whitney, control vs. slurry,  $p = 0.010$ ). Generally, the  $5\beta$ -stanol concentrations in the soil cores decreased exponentially with distance downslope, with average concentrations at the bottom of the slope of  $84\text{ ng g}^{-1}$  which were not significantly different from the control soil (Mann–Whitney, control vs. slurry,  $p = 0.343$ ). The average concentration of the eroded sediment was  $2040\text{ ng g}^{-1}$  which was also significantly enriched compared with the control soil (Mann–Whitney, control vs. slurry,  $p = 0.010$ ). Figure 10 shows the changes in the concentration of  $5\beta$ -stanols in the soil cores in the downslope direction after each individual experiment. Some of the experiments showed increases in the concentrations of soil core  $5\beta$ -stanols at the bottom of the slope, which suggests that deposition of slurry-derived IOM was occurring close to the slope outlet. The error bars in Fig. 10 represent the range of values from the three soil cores taken horizontally at each distance downslope. In some locations the range of concentrations was large, possibly due to preferential surface runoff pathways which distributed sediment laterally across the slope as well as downslope.

In addition to absolute concentrations the ratio between  $5\beta$ - (slurry-derived) and  $5\alpha$ - (soil-derived) stanols was calculated in order to assess the contribution of slurry-derived IOM vs. native soil IOM in the soil cores (data not shown). Results showed

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a spatial pattern the distribution of the ratio of  $5\beta$ - :  $5\alpha$ -stanols in the soil across the slope surface for each experiment, exhibiting an exponential decline in  $5\beta$ - :  $5\alpha$ -stanols ratio in the downslope direction from where slurry was applied. The main difference between the experiments was in the variability of stanol ratios, which was a function of slope gradient. The standard deviation of the stanol ratios was calculated for each experiment (not including the top of the slope where the slurry was initially applied) in order to assess the variation in transported slurry-derived material between experiments. The analysis showed that increasing slope gradient resulted in an increase in the standard deviation of the ratios in soil cores across the slope suggesting that surface runoff at lower gradients was flowing more evenly over the soil surface compared to the steeper slopes which may have exhibited more preferential flow routing.

The concentrations of 5 $\beta$ -stanols were also quantified in the sediment eroded from the slope during each experiment. In order to obtain an adequate sample of sediment for 5 $\beta$ -stanol analysis, the eroded sediment samples were combined into three time segments, representing the beginning, middle and end of the rainfall simulation (see Table 1 for the experimental time covered by each composite sample). As the sediment flux from the 5°, 60 mm h<sup>-1</sup> experiment was relatively low only one analysis was possible which represented the entire event. The average concentrations of 5 $\beta$ -stanols are shown in Fig. 9 and the temporal variations in concentrations are shown in Table 2. The lowest concentration of 5 $\beta$ -stanols was recorded in the eroded sediment from the experiment with the lowest volume of surface runoff, suggesting that transport of slurry-derived POM was therefore limited, and therefore reflected in the low sediment yield under these conditions. Experiments with higher volumes of surface runoff resulted in an increase in the 5 $\beta$ -stanol concentrations by up to a factor of 10. Increasing rainfall intensity appeared to have little effect on the concentrations of 5 $\beta$ -stanols exported at the beginning of the rainfall simulations at a slope gradient of 5°. However, increasing slope gradient resulted in a rapid decrease in concentrations during the middle and end of the simulations. The supply of slurry-derived material was exported from the slope faster at steeper slope gradients due to the increased sediment flux, as shown

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in Table 2. Slope gradient had a larger impact on the total transport of 5 $\beta$ -stanols than the increase in rainfall intensity, due to an exponential increase in sediment yield.

## 4 Discussion

This study used a series of six controlled laboratory experiments to assess the effect of slope gradient and rainfall intensity on the transport of slurry-derived compounds. This work also provided an extension to the proof of concept developed in Lloyd et al. (2012) for using fluorescence and 5 $\beta$ -stanols simultaneously to trace both the vertical and lateral transport of slurry-derived material in both dissolved and particulate forms.

#### 4.1 Role of slope gradient and rainfall intensity on transport slurry-derived compounds

It was hypothesised that changes in slope gradient and rainfall intensity would alter the relative partitioning of slope flow pathways and erosion rates which would in turn exert a strong control on the flux and yield of slurry-derived compounds. The impact of these variables on slurry transport can be described by two main hydrologically-driven scenarios.

(1) *Overland flow dominated systems, which typically occur on steep slopes and/or high rainfall intensities.* High overland flow rates typically result in high erosion rates and the combination of the two enhances transport of slurry-derived compounds via surface pathways (dissolved in runoff and bound to eroded sediment), which are rapid and interact directly with the source of applied slurry material. The concentration of the dissolved components depends on the volume of runoff produced. Higher intensity rainfall events generally transported the overall largest load of slurry-derived dissolved material but the concentrations were lower. This result is supported by Delpla et al. (2011), who showed in a field study that it was the highest intensity rainfall events which generated overland flow that yielded the highest DOC export prior to cattle slurry

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application. Dissolved components of slurry which have been diluted by increased volumes of discharge will have a lower immediate impact on stream ecology. However, the larger total load exported during high intensity storms will result greater total losses of nutrients from the soil and could result in pressures on receiving waters, such as increased biological oxygen demand (BOD). It is estimated that input of cattle slurry can produce a BOD of between 10 000 and 30 000 mg L<sup>-1</sup>, leading to reduced oxygen levels and ultimately the death of aquatic life (Khaleel et al., 1980; Baker, 2002b; Foy and Kirk, 1995).

The concentrations of slurry-derived particulates were controlled primarily by the slope gradient, however, the overall load increased substantially (between 1 and 3 orders of magnitude) with increased rainfall intensity due to the increased sediment transport by overland flow. This is supported by Michaelides et al. (2010) who showed the importance of erosion events on landscape nutrient loss. Although small amounts of subsurface and percolated flow occurred at the higher rainfall intensity, they do not play a key role in the transport of slurry-derived particulates. In the overland flow dominated experiment (10°, 120 mm h<sup>-1</sup>) over 4 kg of sediment were exported (over an area of 15 m<sup>2</sup>), containing 2.3 mg of 5β-stanols in under an hour of rainfall, compared with a total of ~9.7 mg of 5β-stanols added to the soil as slurry before the experiment. The experiment that generated less than 1 % of its discharge by overland flow (5°, 60 mm h<sup>-1</sup>) eroded just 14 g of sediment (3 μg 5β-stanols) during the experiment. Due to the rapid movement of slurry-derived material in this overland flow scenario there is little time for the slurry derived components to transform, for example the nitrification of NH<sub>4</sub><sup>+</sup> to NO<sub>2</sub><sup>-</sup> and NO<sub>3</sub><sup>-</sup> or the mineralisation of the organic fraction to more labile inorganic forms. As a result, over short time scales after slurry application, the transported fractions of slurry tended to reflect the original slurry composition.

(2) *A predominance of vertical, percolated flow, which typically occur due to low slope gradients coupled with lower rainfall intensities.* Under this regime, the export of slurry-derived material is mainly in dissolved form and the flux tends to be higher directly beneath, or close to the application area. The timing of the dissolved flux also tends to



allowing diffusion to liberate chemicals from the soil matrix. Experiments run by Gao et al. (2004) showed that at the beginning of a rain event raindrop impact was the main mechanism for liberating solute, followed by diffusion at the latter stages of the storm. In addition, research has also shown that storms which have variable rainfall intensities can increase the transport of solute in surface runoff and via preferential subsurface routes (Zhang et al., 1997; Malone et al., 2004).

4.2 Use of a combined tracer approach to monitor the transport of slurry-derived material.

This study also aimed to test efficacy of the use of a combined biomarker approach to monitor the vertical and lateral transport of dissolved and particulate slurry fractions. In the current study the transport of dissolved slurry-derived material within different flow pathways was monitored using NH<sub>4</sub><sup>+</sup> concentrations and fluorescence. Over short time-scales (hours) NH<sub>4</sub><sup>+</sup> concentrations can provide useful information regarding the timing and spatial patterns of movement of dissolved slurry compounds. Field data have shown that areas which have high N-loading, such as those treated with slurry, could receive N at a higher rate than can be incorporated into the organic fraction, resulting in inorganic-N, especially NH<sub>4</sub><sup>+</sup> becoming available for rapid transfer to water-courses, often via preferential flow pathways (Heathwaite and Johnes, 1996). However, within days the signal will degrade due to nitrification of NH<sub>4</sub><sup>+</sup> into NO<sub>3</sub><sup>-</sup>, which would be difficult to distinguish from soil-derived N, without the use of isotopic enrichment additions.

Fluorescence spectroscopy was shown by Lloyd et al. (2012) to be a robust tracer of DOM, which was slurry-derived over short timescales and in relatively small-scale soil columns. However when this technique was applied to the more complex 3-D laboratory system the results were not as clear. While the fluorescence intensity at an excitation at 270 nm was not significantly different between the slurry-treated and control experiments, using the ratio between “270” and “360” intensities provided a detectable signal. The ratio was able to identify smaller changes in the fluorescence spectra which

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were not possible using the “270” fluorescence intensity alone. The fluorescence intensities recorded from the TRACE experiments were several orders of magnitude lower than those from the soil column experiments (Lloyd et al., 2012), due to the larger volume of rainfall added in the TRACE experiments which resulted in a significantly smaller slurry: soil ratio. The slurry was only applied to the top 1 m strip of the slope and therefore the signal was rapidly diluted both by the rainfall and the additional signal from the soil-derived DOM. Furthermore, the “270”: “360” ratio of the slurry was calculated at 1.7, which is lower than the range cited in the literature ( $\sim 2$ –5) for cattle slurries (Baker, 2002b). This could be because when the slurry was collected (during April 2008), it had been a very wet spring and as a result the farm slurry store had received higher than average volumes of rainwater, potentially diluting the slurry before application.

Fluorescence spectroscopy has been shown previously to be a powerful technique for characterising DOM, even in very dilute samples (e.g. Barker et al., 2009; Birdwell and Engel, 2010). However, the signal was more difficult to detect in this case because the soil-derived DOM swamped the slurry signal, making it difficult to resolve the two sources. This problem was exaggerated in the case of the higher intensity rainfall experiments due to the additional water being transported with the slurry. If a larger section of the slope had been treated with slurry then it is more likely that the signal could have been seen. Naden et al. (2010) showed that drainage water from field lysimeters treated with slurry could be distinguished from control experiments, even though only a small proportion of the applied slurry was thought to have been leached from the system. However, work by Old et al. (2012) showed that fluorescence differences could not be identified in storm runoff samples after an application of cattle slurry to undrained plots. The authors suggest that this was due to rapid absorption or microbially-mediated-immobilisation of the slurry material in the soil matrix. Baker (2002a) showed that river water samples from the UK exhibited “270”: “360” ratios of  $0.37 \pm 0.41$  ( $n = 242$ ), suggesting that cattle slurries may be difficult to detect once they are diluted compared to the background river fluorescence. However, Baker (2002a)



conclude that the techniques would be able to detect cattle slurry inputs from large point sources, such as slurry tank failures.

The particulate fraction was investigated using bulk elemental analysis, followed by quantifying the  $5\beta$ -stanol concentrations. Soil analyses of bulk C and N were found to be generally ineffective at tracing transport pathways of slurry-derived POM. The eroded sediments from all of the experiments showed increases in TOC and TN values compared with the control but the source of origin of the TOC and TN cannot be identified (i.e. slurry or soil). The use of the specific biomarkers  $5\beta$ -stanols provides more robust information about the transport of slurry-derived particulates as they have been shown to be an unequivocal tracer of ruminant slurry (Bull et al., 2002; Elhmmali et al., 2000; Evershed et al., 1997; Nash et al., 2005). The concentration of the  $5\beta$ -stigmastanol in the slurry used in this study was lower than other published values, with an average concentration of  $\sim 12 \mu\text{g g}^{-1}$  of freeze dried slurry, compared with  $\sim 46 \mu\text{g g}^{-1}$  (Leeming et al., 1996). However, there was still a large difference between the slurry concentration and the initial soil, so this did not cause any issues for using the biomarker as a tracer. On the other hand, the ratios of the  $5\beta$  :  $5\alpha$ -stanols determined in the current study were very similar to other published values, where ratios from soil cores taken from the slurry application area ranged from 2.4–2.9, compared with an average ratio of  $\sim 2$  measured in cow manures (Evershed et al., 1997). Data collected here further illustrate that  $5\beta$ -stanols are a robust tracer of slurry-derived POM, and therefore could be used to monitor the transport of particulate forms of slurry.

Overall, the results show that using a combination of tracers and biomarkers for both soluble and insoluble fractions can be very effective for tracing the movement of slurry via multiple transport pathways. This work provides a proof of concept that the methodology works in larger and more complex controlled laboratory systems. While elements of this type of methodology have been tested in field scenarios (e.g. Naden et al., 2010; Granger et al., 2010; Evershed et al., 1997; Bull et al., 1998; Nash et al., 2005), there is still a need to test this combined tracer approach in a larger field study.

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This work has provided important and new insights into flow partitioning across a range of controlled hillslope and rainfall scenarios and has allowed quantification of the impact this has on the transport of slurry-derived compounds. Results indicate that the dissolved components of slurry-derived material (traced using  $\text{NH}_4^+$  and the ratio between “270” : “360” fluorescence intensities) were transported rapidly through the soil system, while the predominant pathway depended on the flow partitioning. When the conditions favoured surface runoff, i.e. high slope gradients and/or high rainfall intensities, larger quantities of slurry-derived material were moved in the surface and sub-surface flow pathways. These shallow flow pathways transport water more rapidly to the slope outlet compared with vertical percolated flow which travelled slowly through the soil matrix. The movement of slurry-derived particulates (traced using  $5\beta$ -stanols) is driven exclusively by the erosion rates on the slope.

Rainfall events which produced flashy hydrological responses, resulting in large quantities of surface runoff, were likely to move sediment and also flush dissolved components of slurry-derived material from the slope, increasing the contamination risk. Rainfall events which produced slower hydrological responses were dominated by vertical percolated flows removing less sediment-associated material, but produced leachate which could contaminate deeper soil layers, and potentially groundwater, over a more prolonged period.

This work has also provided one of the first examples of using multiple biomarkers to assess the effects of slope gradient and rainfall intensity on the movement of slurry derived OM. The results have shown that this approach can be successfully applied to more complex 3-D systems (than simple soil columns) and can yield valuable data about the interactions between slurry and the soil-water system. Overall, this research provides new insights into the partitioning of slurry-derived material when applied to an unvegetated slope and the transport mechanisms by which contamination risks are created.

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# Runoff- and erosion-driven transport of cattle slurry

C. E. M. Lloyd

**Table 1.** Chemical characteristics of control soil and cattle slurry (dry weight basis) used in all of the experiments.

Parameter	Initial soil	Cattle slurry
Total carbon	2.7 % $\pm$ 0.2	39.6 % $\pm$ 0.3
Total organic carbon	2.6 % $\pm$ 0.2	39.0 % $\pm$ 0.3
Total nitrogen	0.4 % $\pm$ 0.09	3.4 9 % $\pm$ 0.004
Ammonium	0.0016 % $\pm$ 0.0005	0.39 % $\pm$ 0.0005
Nitrate	0.0022 % $\pm$ 0.00006	0 %
Nitrite	0.00004 % $\pm$ 0.00003	0 %
Organic N	0.4 % $\pm$ 0.09	3.45 % $\pm$ 0.004

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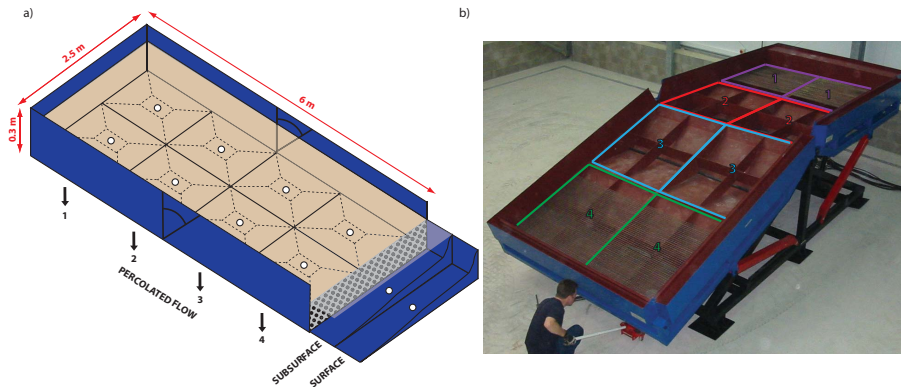
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**Table 2.** Concentrations of  $5\beta$ -stanols in the eroded sediment (dry weight basis) through time and the total exported mass throughout the experiment.

		60 mm h <sup>-1</sup>				120 mm h <sup>-1</sup>				
		5°		10°		5°		10°		
	Time (min)	Conc. (ng g <sup>-1</sup> )	Mass (μg)	Conc. (ng g <sup>-1</sup> )	Mass (μg)	Time (min)	Conc. (ng g <sup>-1</sup> )	Mass (μg)	Conc. (ng g <sup>-1</sup> )	Mass (μg)
Beginning	0–30	351*	3.3	3477.2	666.7	0–20	306.1	3611.0	2024.4	6912.5
Middle	30–60	–	–	3005.2	716.5	20–35	545.0	1032.2	1426.0	9274.2
End	60–100	–	–	2533.1	734.0	35–50	360.7	417.1	1280.2	6725.3
Total			3.3		2123			5060		22 912

\* This experiment only had one time interval due to the lack of sediment to analyse.

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**Figure 1. (a)** Schematic showing sampling outlets on the TRACE slope and **(b)** photo showing the empty slope.

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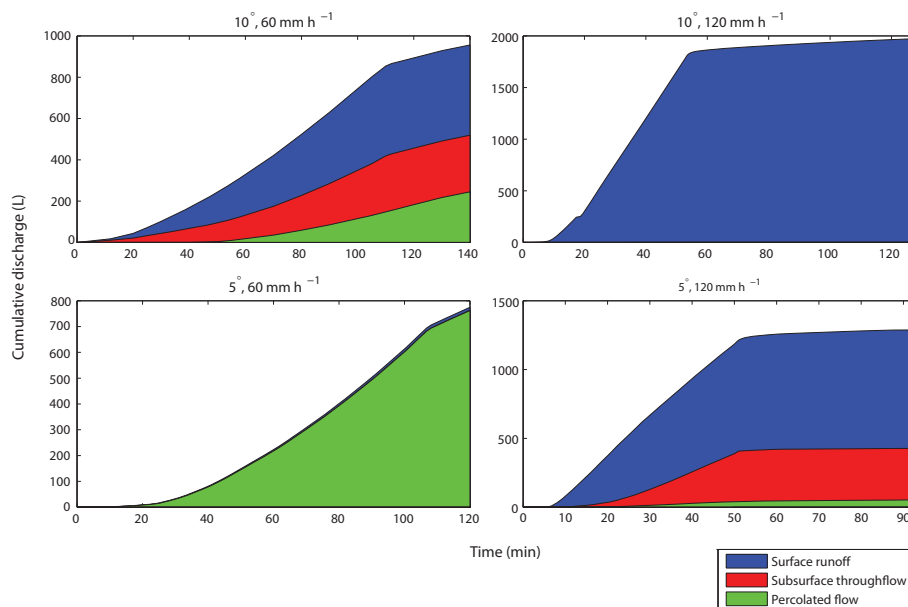
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**Figure 2.** Plots showing cumulative discharge from surface runoff, subsurface throughflow and vertical percolated flow from the slurry-treated slope experiments.

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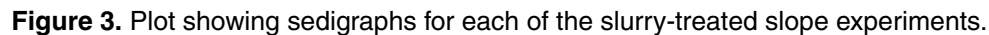
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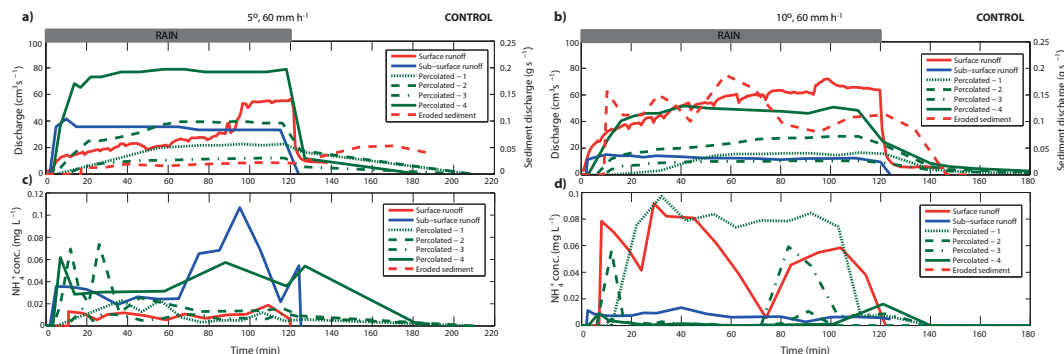
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**Figure 4.** Plots showing water and sediment discharge (**a, b**) and NH<sub>4</sub><sup>+</sup> concentrations (**c, d**) from each monitored flow pathway for the control experiments (no slurry) at two slope angles (5 and 10°) at 60 mm h<sup>-1</sup> rainfall intensity. Where percolated 1–4 represent vertical transport from each of the pairs of tanks beneath the soil slope as marked in Fig. 1.

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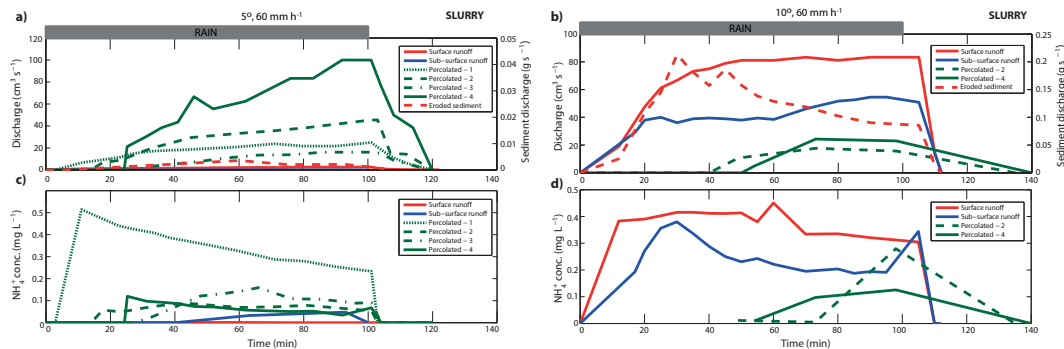
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# Runoff- and erosion-driven transport of cattle slurry

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**Figure 5.** Plots showing water and sediment discharge (**a, b**) and NH<sub>4</sub><sup>+</sup> concentrations (**c, d**) from each monitored flow pathway for the slurry treated experiments at two slope angles (5 and 10°) at 60 mm h<sup>-1</sup> rainfall intensity. Where percolated 1–4 represent vertical transport from each of the pairs of tanks beneath the soil slope as marked in Fig. 1. Note: data which is absent reflects the lack of activation of a particular flow pathway.

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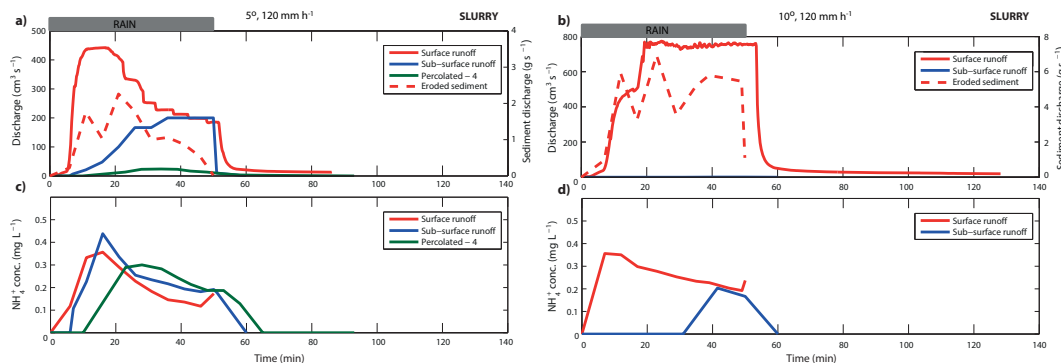
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**Figure 6.** Plots showing water and sediment discharge (**a, b**) and  $\text{NH}_4^+$  concentrations (**c, d**) from each monitored flow pathway for the slurry treated experiments at two slope angles (5 and 10°) at  $120 \text{ mm h}^{-1}$  rainfall intensity. Where percolated 1–4 represent vertical transport from each of the pairs of tanks beneath the soil slope as marked in Fig. 1. Note: data which is absent reflects the lack of activation of a particular flow pathway.

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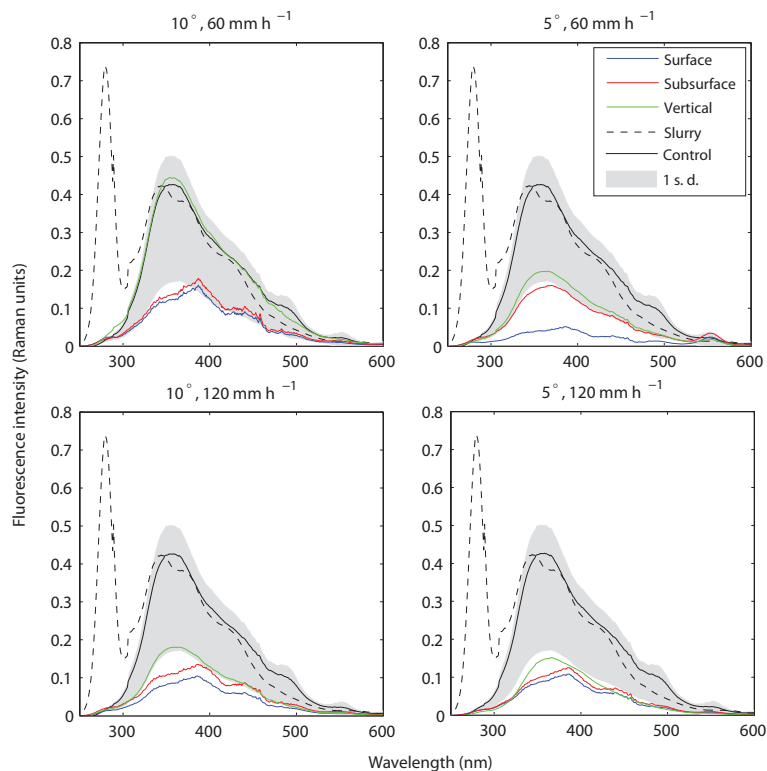
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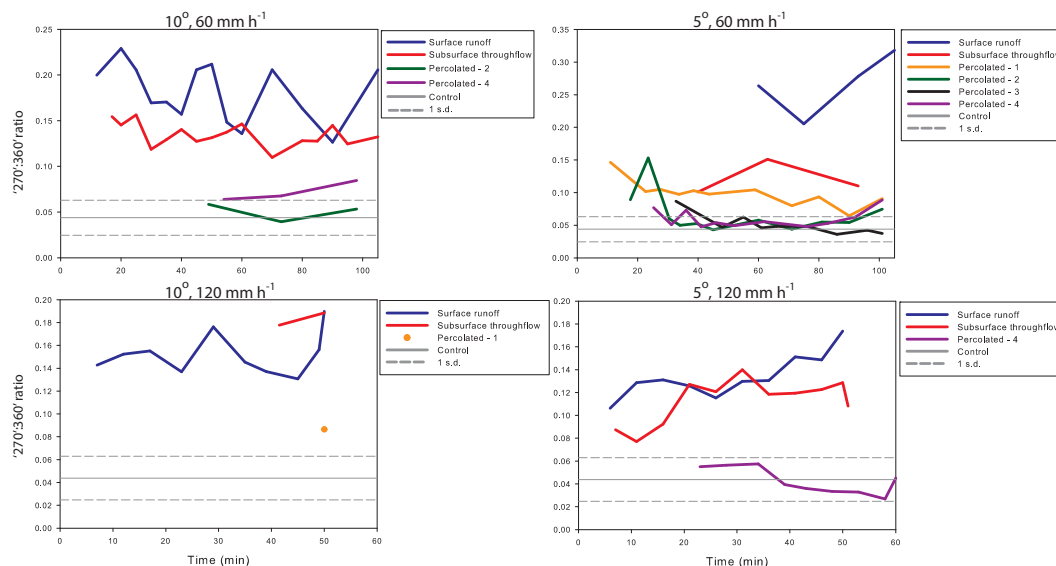


**Figure 7.** Plots showing the mean fluorescence spectra from the surface runoff, subsurface throughflow and vertical percolated pathways for each slurry-treated experiment along with the average spectra from all pathways of the control experiments and the spectra from the applied slurry. The shaded area represents 1 standard deviation around the mean of the control soil spectra.



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**Figure 8.** Plots showing the ratio of “270” : “360” nm fluorescence intensities for each measured flow pathway. The grey lines represent the average value for the control samples and the grey dashed lines are 1 standard deviation around the mean.

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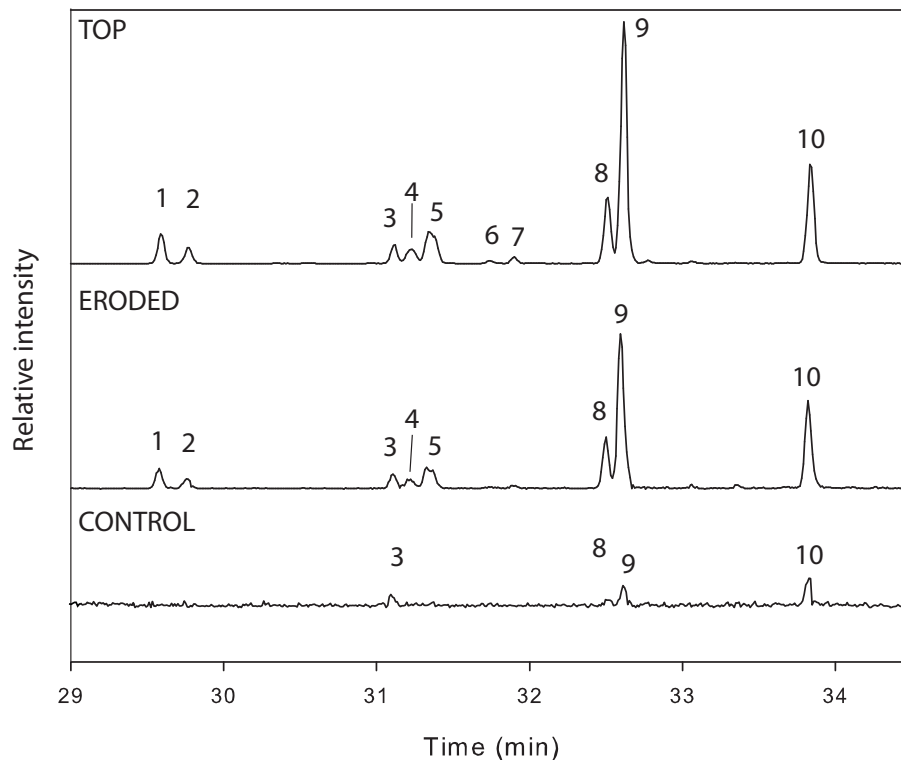
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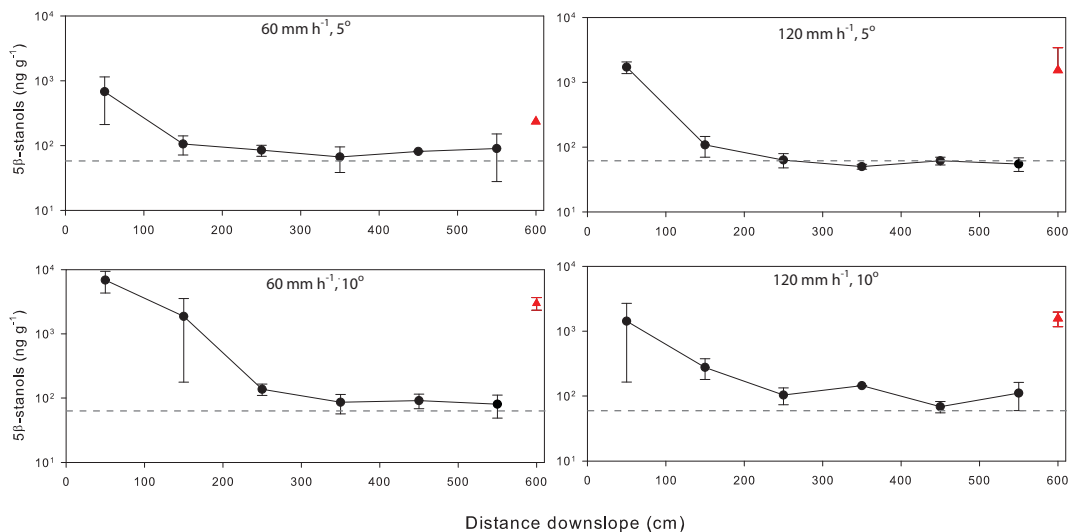




**Figure 9.** Partial chromatogram ( $m/z = 215$ ) showing examples of characteristic hydrolysed lipid extracts from the top section of the soil slope, the eroded sediment (taken from the  $10^\circ$ ,  $60 \text{ mm h}^{-1}$  experiment) and the control soil. Where: 1 = coprostanol, 2 = epicoprostanol, 3 =  $5\alpha$ -cholestanol, 4 =  $5\beta$ -campestanol, 5 =  $5\beta$ -epicampestanol, 6 = 24-ethyl-campestanol, 7 = 24-ethyl- $5\beta$ -cholestan- $3\alpha$ -ol, 8 =  $5\beta$ -stigmastanol, 9 =  $5\beta$ -epistigmastanol, 10 =  $5\alpha$ -stigmastanol.

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**Figure 10.** Plots showing the change in average concentration of  $5\beta$ -stanols in soil cores downslope for each of the slurry-treated experiments. The dashed line shows the mean concentration in the control soil, and the triangles represent the average concentrations in the eroded sediment.

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